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### (54) System and method for waveform morphology comparison

Verfahren und System zum Vergleichen der Morphologie von Wellenformen

Procédé et système pour la comparaison de la morphologie de formes d'ondes

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(56) References cited:  
**WO-A-96/25096** **US-A- 4 589 420**  
**US-A- 5 000 189** **US-A- 5 240 009**

- **GREENHUT S.E. ET AL:** "Separation of ventricular tachycardia from sinus rhythm using a practical, real-time template matching computer system" PACE, vol. 15, no. 2, November 1992, pages 2146-2153, XP002098299

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**Description**

- [0001] The present invention relates generally to comparing waveforms without requiring sample-by-sample comparison. More particularly, the present invention relates to electrogram waveform comparison for discriminating ventricular tachycardia (VT) from sinus tachycardia (ST) and supraventricular tachycardia (SVT).  
 5 [0002] In a heart monitoring system, it is often desirable to distinguish between ventricular complexes that are conducted from the atria (i.e., normal complexes) and ventricular complexes that originate in the ventricular myocardium (i.e., abnormal complexes). For example, it is important to be able to distinguish sinus tachycardia (ST) and supraventricular tachycardia (SVT) from ventricular tachycardia (VT). Waveform morphology comparison is normally used to accomplish such discrimination. That is, a test intraventricular electrogram (IVEG) is compared to a template IVEG. The template IVEG is generally taken during a sinus rhythm. If the morphology comparison indicates that there is a ventricular tachycardia, corrective action is taken. For example, implantable cardioverter defibrillators (ICDs) or pacemakers can undertake corrective action when they detect ventricular tachycardia. For a more detailed description of the origin and morphology of these signals, see US-A-5 240 009 to Williams, entitled "Medical Device with Morphology Discrimination."  
 10 [0003] From this US-A-5 240 009, a medical device is known for monitoring waveform complexes of intracardiac electrograms. An intracardiac complex is said to occur when the signal exceeds an upper threshold. Lower threshold peaks which are part of that complex can occur before and after the upper threshold crossing. For each complex, there is a time limit and a limit on the number of peaks allowed. Thus the maximum number of peaks within a complex is limited by a given number, for example five. A standard complex is computed by averaging identification criteria (amplitude, width and plurality of peaks) of several complexes. Subsequent complexes are then compared to the stored standard complex. Such comparison includes comparing peaks of subsequent complexes with the peaks of the stored standard complex, aligning subsequent complexes with the stored standard complex, and providing a score associated with the comparisons and alignment.  
 15 [0004] A major consideration in developing ICDs is their limited battery power. After implantation, batteries cannot be replaced without surgery and generally the entire ICD is replaced when the batteries are depleted. Thus, it is desirable to conserve battery power. One way to conserve battery power is by reducing the complexity of the signal processing that must be performed by the ICD.  
 20 [0005] Conventional morphology algorithms compare test and template IVEGs on a point-by-point basis. That is, characteristics about the points in the test IVEG are compared to corresponding points in the template IVEG. Examples of such point-by-point morphology matching processes are disclosed in Greenhut, Saul E. & Steinhaus, Bruce M., *Template Matching Techniques of Electrophysiologic Signals: A Practical Real-Time System for Detection of Ventricular Tachycardia*, Biomedical Scientific Instruments, 1992, Vol. 28, pp. 37-42 (hereinafter "Template Matching Technique") and Greenhut, Saul E. et al., *Separation of Ventricular Tachycardia From Sinus Rhythm Using a Practical, Real-Time Template Matching Computer System*, PACE 1992, 15:2146-2153 (hereinafter "Separation of Ventricular Tachycardia").  
 25 [0006] Such conventional waveform comparison techniques suffer from several shortcomings. First, waveform alignment is critical to a proper point-by-point comparison. If the test and template signals are not aligned correctly, the result of the waveform comparison can be erroneous. Furthermore, aligning the test and template signals can be a burdensome and time consuming problem.  
 30 [0007] Second, these conventional point-by-point processing techniques are computationally intensive. For example, such techniques generally require two divide operations. This is illustrated in the equation for the normalized area difference (NAD) in the references cited above (see equation (1) on page 39 of "Template Matching Techniques" and equation (1) on page 2148 of "Separation of Ventricular Tachycardia").  
 35 [0008] Both of these disadvantages result in excess current drain on the battery of an implanted ICD. The excess current drain reduces the life expectancy of the ICD. The reduced life expectancy of the ICD exposes the patient to a greater risk and expense of repeat surgery to replace the device.  
 40 [0009] Thus, what is needed is an improved method for discriminating ventricular tachycardia (VT) from sinus tachycardia (ST) and supraventricular tachycardia (SVT). Such a method should reduce computational complexity for power savings while at the same time reducing the alignment problem.  
 45 [0010] The present invention is directed to distinguishing between a first signal and a second signal on the basis of how well the morphologies of the first and second signals match. The present invention forms a template based on the morphology of a "known" signal. The "known" signal can be, for example, a signal collected during a period where a patient is known to exhibit a normal sinus rhythm. A test signal is compared against the template to determine how closely the signals correspond using their morphology.  
 50 [0011] In the preferred embodiment, the morphology is represented by a consecutive series of N data samples. From these N data samples, M (e.g., 3) peaks are identified. The positions (i.e., order of occurrence), polarities and normalized areas of the peaks are compared to align the signals and yield a score. A higher score indicates less similarity.

between the first and second signals.

[0012] In the preferred embodiment, the invention is applied to monitoring heart signals to analyze the morphology of an intraventricular electrogram (IVEG) for the purpose of discriminating ventricular tachycardia (VT) from sinus tachycardia (ST) and supra ventricular tachycardia (SVT). A template is generated by analyzing an IVEG depolarization during a sinus rhythm. The polarity and normalized area of the most significant consecutive peaks of the IVEG depolarization are recorded as the template. Each sensed IVEG is matched against the template by comparing the position, polarity and normalized area of the most significant consecutive peaks and summing the differences. The resulting value provides a measure of how similar in morphology the sensed IVEG is to the template IVEG.

[0013] Further features and advantages of the invention, as well as the structure and operation of various embodiments of the invention, are described in detail below with reference to the accompanying drawings. In the drawings, like reference numbers generally indicate identical, functionally similar, and/or structurally similar elements. The drawing in which an element first appears is indicated by the digit(s) to the left of the two rightmost digits in the corresponding reference number.

[0014] The present invention will be described with reference to the accompanying drawings, wherein:

FIG. 1 is a flow chart for a method of morphology comparison of two signals according to a preferred embodiment.  
 FIG. 2 is a graphic representation of an IVEG collected during a morphology window.

FIG. 3 is a flow chart for determining peak information of interest according to the preferred embodiment.

FIG. 4 is a flow chart for aligning a test and a template IVEG according to a preferred embodiment.

FIG. 5 is a flow chart for determining whether peaks match according to a preferred embodiment.

FIG. 6 is a flow chart for determining whether peak positions match and for correcting mismatch according to a preferred embodiment.

FIG. 7 illustrates an example mismatch correction using the flow chart of FIG. 6.

FIG. 8 is a system for morphology comparison of two signals according to a preferred embodiment.

FIG. 9 is a schematic representation of a preferred implementation of the system for morphology comparison of two signals in FIG. 8.

[0015] The template matching morphology method of the present invention calculates the difference in the normalized area of each significant peak of a template electrogram (EGM) and a test EGM over a fixed window size of data samples. In addition, the test and template EGMs are not limited to collection from a patient. For example, the test and template EGMs can be computer generated for use, for example, in technique testing.

[0016] For purposes of illustrating a preferred embodiment, the invention is described in terms of comparing intraventricular electrograms (IVEGs). It should be understood, however, that this is done for purposes of illustration only. The invention can be used for comparing any cardiac signal whether it is an EGM taken from implanted electrodes or an electrocardiogram (ECG) from surface electrodes or otherwise.

[0017] Using a device designed according to the preferred embodiment of the present invention reduces the complexity of waveform morphology diagnosis. As a result, power is conserved and battery life is extended.

[0018] A flow chart for a template morphology matching method 100 according to a preferred embodiment of the invention is described with reference to FIG. 1. The beginning of method 100 is indicated in a step 102. In a step 104, a template IVEG is created. The template IVEG is analogous to a standard to which test signals are compared. The template can be generated in a number of well known ways. In the preferred embodiment, the template is generated by simply collecting data points (i.e., digital samples of the IVEG) during a morphology window for a complex in a sinus rhythm. In an alternate embodiment, data is collected over a plurality of morphology windows during which there is a sinus rhythm. After the data is collected, the data is averaged to determine a template. Prior to the averaging however, the data must be aligned. A data alignment mechanism such as described herein can be employed.

[0019] As part of the template creation step, the IVEG data is normalized. Normalization accounts for differences in absolute magnitude of the sensed signals that may be affected by variables such as electrode placement, differences in amplifier gain, and the like. Normalization allows the method of the invention to compare the relative magnitudes of the peaks in the IVEG. Normalization of the template IVEG is described in detail below.

[0020] The template is generated using morphology information of interest in the signal. In the preferred embodiment, the morphology information selected for comparison is information about M (e.g., 3) consecutive peaks. The peak information includes normalized area, polarity, and position (i.e., order of occurrence) for the M consecutive peaks having the largest sum of peak amplitudes in a collected signal. In the preferred embodiment, the signals are IVEG depolarizations.

[0021] After creation of the template IVEG, a test IVEG is collected in a step 106. As discussed below with reference to FIG. 2, the test IVEG is collected over a morphology window 201 in a well known manner.

[0022] Next, in a step 108, peak information is identified in the test IVEG. The test and template IVEGs are then aligned in a step 109. Alignment of the test and template IVEG's is tested in a step 110. If alignment failed, then method

100 proceeds to step 114 where the test and template IVEGs are indicated as being dissimilar. If alignment was successful (as indicated at step 110), then method 100 proceeds to step 111.

[0023] In step 111, the test and template IVEGs are compared to determine a score. The score represents how well the test and template IVEGs match. The lower the score, the more similar are the test and template IVEGs. The score is compared to a threshold in a step 112. If the score does not exceed the threshold, then the test and template IVEGs are indicated as being similar in a step 116. If the score exceeds the threshold, then the test and template IVEGs are indicated as being dissimilar in step 114.

[0024] In the preferred embodiment, the threshold of step 112 is determined by looking at a number of trial runs of a patient in a sinus rhythm and a VT rhythm. The scores are then plotted and viewed. A threshold value is picked that will provide an acceptable (normally very low) probability that a score indicating a VT rhythm is interpreted as a score for a normal sinus rhythm and vice versa.

[0025] After either of steps 114 or 116 are executed, method 100 ends as indicated at step 118. Alternatively, method 100 may return to step 106 and repeat. In this manner, method 100 can continuously monitor an input stream of test IVEGs.

[0026] As explained in further detail below, this method of the present invention significantly reduces computational complexity. Because the method calculates a normalized area of peaks, there are optimally only three divides required. Importantly, the method of the invention improves over conventional methods by not requiring two divides per sample point. Moreover, the alignment problem is solved by aligning on the basis of morphology (i.e., peak polarity and peak amplitude), rather than corresponding data points (samples).

[0027] The steps of method 100 are now explained in further detail with reference to FIGS. 2-8.

[0028] FIG. 2 illustrates a graphical representation of a single IVEG depolarization to be collected in step 106. The test IVEG is collected and sampled during a morphology window 201 in a well known manner. Morphology window 201 begins at a time 202 and ends at a time 204. During the morphology window 201, N data samples of the IVEG depolarization are taken.

[0029] Referring still to FIG. 2, step 108 of determining peak information of the test IVEG is described. A peak is defined to begin with the data sample that is greater than (in absolute magnitude) or equal to a fixed threshold 206 or 208 (in absolute magnitude). A peak is defined to end with the data sample that crosses the baseline 207. Three peaks 210, 212 and 214 are illustrated. Peak 212 is the dominant peak, peak 214 is the second dominant peak, and peak 210 is the third dominant peak. The peaks must be consecutive.

[0030] FIG. 2 also illustrates a sensing window 218. Sensing widow 218, also known as the sense refractory period, begins when a peak rises above a sensing threshold 216. The end of the sensing window 218 corresponds to the end of morphology window 201.

[0031] As illustrated in FIG. 2, peaks may be collected both before and after sensing window 218 begins. In the present invention, M (e.g., 3) peaks having the largest sum of peak amplitudes are used to analyze the waveform morphology. While three peaks are preferred in the present implementation of the invention, a person skilled in the art will recognize that a different number of peaks (e.g., 2, 4 or more) may be used.

[0032] FIG. 3 illustrates in further detail step 108 (see FIG. 1) of determining peak information for the test IVEG. The peak information includes the normalized area of each peak, the polarity of each peak, and the positions (not in absolute time, but in order of occurrence) of the dominant and second dominant peaks in the test IVEG.

[0033] When performing step 108, the method of peak determination begins in a step 302. In a step 303, the three consecutive peaks with the largest sum of peak amplitudes are extracted. Next, in a step 304, the area is calculated for each of the three peaks selected in step 303. The area is essentially the size of the peak. In the preferred embodiment, the area is simply the arithmetic sum of the data points (amplitudes) in the peak. It would be apparent to those skilled in the art that other methods for calculating area can be used.

[0034] After determining the area for each of the three peaks selected in step 303, the method executes step 306. In step 306, the polarity of each of the peaks selected in step 303 is determined. Polarity refers to whether a peak is greater than zero (positive) or less than zero (negative). For example, peak 212 has positive polarity, whereas peaks 210 and 214 have negative polarities.

[0035] After determining the polarity for each of the peaks, the positions of the most dominant and second most dominant peaks in the three-peak set are determined in a step 308. Note that, in the preferred embodiment, peak dominance is measured by peak amplitude, rather than by peak area. Also note that this position determination refers to a relative position or order of occurrence (e.g., third dominant peak in a first position, dominant peak in a second position, and second dominant peak in a third position as for Fig. 2) rather than an absolute position (e.g., dominant peak at 63 msec, second peak at 71 msec, etc.) within the morphology window.

[0036] After determining the relevant positions, the area of each peak is normalized in a step 310. In the preferred embodiment, the normalization proceeds according to Equation (1) for each peak in the 3-peak set selected:

## Equation (1)

$$5 \quad NormAreaPeak_x = \left| \frac{AreaPeak_x}{|AreaPeak_1| + |AreaPeak_2| + |AreaPeak_3|} \right|$$

In an alternate preferred embodiment, the normalized area of each peak is calculated according to Equation (2).

10

## Equation (2)

$$NormAreaPeak_x = \left| \frac{AreaPeak_x}{|PeakWithMaxArea|} \right|$$

15 The subscript  $x$  in Equations (1) and (2) indicates that the calculation is done for each peak selected in step 303. After normalization, the method ends at step 312 to complete the determination of peak information for the test IVEG.

[0037] Step 109 of FIG. 1 represents alignment of the test and template IVEGs. This step is detailed in FIG. 4. The method of FIG. 4 begins in a step 402. In a step 403, an attempt is made to match the polarity of a dominant peak in the template IVEG with the polarity of a dominant peak in the test IVEG. Step 403 is described in further detail below with reference to FIG. 5.

[0038] A step 404 determines whether the peak polarity matching of step 403 was successful. If decision step 404 determines that peak polarities do not match, then step 407 determines that the alignment failed and the method continues to step 408. In step 408, the method of step 109 ends and returns to the method of FIG. 1. If, on the other hand, step 404 determines that the peak polarities do match, then the method continues to step 405.

20 [0039] Step 405 determines whether the peaks with matching polarities in the test and template IVEGs correspond to one another in position. Peaks that correspond are peaks having the same sequential position in a particular IVEG. For example, if the peak positions are numbered from one to three, the peak in position one in the test IVEG corresponds to the peak in position one in the template IVEG. If step 405 indicates that the peaks in the test and template IVEGs correspond to one another in position, then the method continues to step 408. In step 408, the method of step 109 ends and returns to the method of FIG. 1.

25 [0040] If, on the other hand, decision step 405 determines that the peaks in the test and template IVEGs do not correspond in position, then the method of step 109 continues with step 406. In step 406, the method attempts to align the peaks by shifting the dominant peak of the test IVEG to the right or the left. The effect of this shifting is to drop a non-dominant peak off of one side of the three peak group and to add another peak to the other side of the three peak group. This will align the peaks between the test and template IVEGs. Step 406 is described in detail below with reference to FIG. 6.

30 [0041] Step 403 of matching peak polarities in the test IVEG with peaks in the template IVEG is now described in further detail with reference to FIG. 5. The method begins in a step 502. In step 504, the polarity of the dominant peak of the template IVEG is compared to the polarity of the dominant peak of the test IVEG. The result of the comparison is checked in a step 506. If the polarities match, then a successful match is indicated at step 508. The method then ends and returns to the method of FIG. 4, as indicated at step 518.

35 [0042] If, at step 506, it is determined that the polarities of the dominant peaks do not match between the test and template IVEGs, then the method continues to step 510. In step 510, the method compares the polarity of the second dominant peak of the template IVEG with the dominant peak of the test IVEG. The method checks the result of the comparison in decision step 512. If the polarities match, then a successful match is indicated at step 508. The method then continues to step 518. If matching is done on the basis of the "second dominant" peak of the template IVEG, this second dominant peak will be called the "dominant" peak of the template IVEG for the remainder of the method. This is done for ease of explanation.

40 [0043] If, at step 512, it is determined that the polarities of the second dominant peak of the template IVEG and the dominant peak of the test IVEG do not match, then the method continues to step 516. In step 516, the method indicates that the peak polarity matching has failed. The method then continues to step 518. If peak polarity matching has failed, then alignment of the test and template IVEGs is not possible. This causes a failure of alignment in step 407 of FIG. 4.

45 [0044] Step 406 of shifting peaks in the test IVEG is now described in greater detail. Step 406 is only reached if the initial peak polarity matching of step 403 (see FIG. 4) is successful, but the positions of the dominant peaks do not correspond (recall that the "second dominant" peak of the template IVEG is considered the "dominant" peak if matching is done on that basis). Step 406 attempts to make the peaks of the test IVEG correspond in position to the peaks of the template IVEG.

55 [0045] Step 406 is described in detail with reference to FIG. 6. The method of FIG. 6 begins at step 602. In step 604,

It is determined whether the position of the dominant peak in the test IVEG is greater than the position of the dominant peak in the template IVEG (recall that the "second dominant" peak of the template IVEG is considered the "dominant" peak if matching is done on that basis). If the test dominant peak has a greater position than the template dominant peak (e.g., test peak at position #2 > template peak at position #1), then it may be possible to align the peaks by shifting the test peaks to the left. A shift does not actually occur. Rather, the test IVEG is inspected or scanned to the right side of the dominant peak. If a new peak is found that matches (in polarity and position) the peak to the right of the dominant peak of the template IVEG, then that new peak is added to the three peak group and the peak that previously occurred to the left of the dominant peak is dropped. Thus, a new group of test peaks is formed. This is indicated by step 605.

[0046] If the test dominant peak has a lesser position than the template dominant peak (e.g., test peak at position #2 < template peak at position #3), then it may be possible to align the peaks by shifting the test peaks to the right. As explained above, a shift does not actually occur. Rather, the test IVEG is inspected or scanned to the left side of the dominant peak. If a new peak is found that matches (in polarity and position) the peak to the left of the dominant peak of the template IVEG, then that new peak is added to the three peak group and the peak that previously occurred to the right of the dominant peak is dropped. Thus, a new group of test peaks is formed. This is indicated by step 606.

[0047] Note that in performing this searching or scanning of steps 605 and 606, the three peaks that are chosen from the test IVEG must be consecutive. Thus, if a mismatching peak occurs on the left side of the dominant peak of the test IVEG, then only the right side can be checked for a matching peak. Similarly, if a mismatching peak occurs on the right side of the dominant peak of the test IVEG, then only the left side can be checked for a matching peak. If a new peak is found in either of steps 605 or 606, the new peak is added to the test IVEG and the mismatched peak is dropped from the test IVEG, as described above, to form a new 3 peak group for the test IVEG.

[0048] After scanning left or right, the method continues in decision step 608 to determine whether the scanning was successful in matching peaks. If no match can be made, the method proceeds to step 614 where the method ends and returns to the method of FIG. 4. If a new peak was found so that the polarities of the three new peaks in the test IVEG match the corresponding peaks in the template IVEG, then the method continues in step 610.

[0049] In step 610, the method selects the three new peaks of the test IVEG for comparison to the template IVEG in step 111 (see FIG. 1). The method then continues to step 612. In step 612, the three new peaks of the test IVEG are normalized using either of Equation (1) or Equation (2) above. The method then continues with step 614. In step 614, the method returns to FIG. 4, and continues in step 408.

[0050] An example of the correction of step 406 is illustrated in FIG. 7. FIG. 7 illustrates a template IVEG 700 and a test IVEG 710. Dashed vertical lines have been drawn over the two IVEGs to show the correspondence therebetween. Note that template IVEG 700 has a dominant peak A, a second dominant peak B, a third dominant peak C and a fourth dominant peak D. The three dominant peaks occur in the order of: C-A-B. Note that test IVEG 710 also has a dominant peak A, a second dominant peak B, a third dominant peak C and a fourth dominant peak D. However, note that the three dominant peaks occur in a different order with respect to the peaks of template IVEG 700. The three dominant peaks of the test IVEG occur in the order of: A-B-C. Thus, step 405 of FIG. 4, would return an indication that the peaks do not correspond in position. This non-correspondence is illustrated in the following table:

IVEG	Position #1	Position #2	Position #3
Template	C	A	B
Test	A	B	C

[0051] Correction of this misalignment according to step 406 of FIG. 4 would proceed as detailed in FIG. 6 as follows. Because peak A (i.e., the dominant peak) of test IVEG 710 occurs first in the three peak group, it is in position #1. However, peak A of template IVEG 700 occurs after peak C. Thus, peak A of template IVEG 700 is in position #2. Step 604 determines whether the test peak A has a greater position than template peak A (i.e., is test peak A at position #1 > template peak A at position #2). In this case, the test peak position is not greater than the template peak position. Therefore, the method proceeds to step 606 where the test IVEG is shifted right (i.e., the IVEG is inspected to the left side of the current three peak group in an attempt to find the next consecutive peak). This operation will locate fourth dominant peak D.

[0052] Note that peak D matches (in polarity and position) (see step 608) peak C of the template IVEG. Therefore, peak D is added to the three peak group and peak C is dropped from the three peak group for the test IVEG. This shift operation results in alignment of the peaks between the test IVEG and template IVEG as illustrated in the following table:

IVEG	Position #1	Position #2	Position #3
Template	C	A	B
Test	D	A	B

5 Note that peak D is taken as the third dominant peak of the test IVEG even though it is really fourth largest in amplitude. This correction to the three peak group of the test IVEG results in all peaks of the test IVEG having matching peaks of the template IVEG.

10 [0053] In the preferred embodiment, the three new peaks of the test IVEG may then be normalized (as indicated in step 612), and a score calculated in accordance with step 111 of FIG. 1. The following equations can be used to calculate a score: where:

15 UnmatchedTemplatePeaks = Normalized Areas of the unmatching template peaks  
 UnmatchedTestPeaks = Normalized Areas of the unmatching test peaks

Equation (3)

20 
$$\text{score} = \sum |\text{UnmatchedTemplatePeaks}| + \sum |\text{UnmatchedTestPeaks}| + \sum |\text{MatchedTemplatePeaks} - \text{MatchedTestPeaks}|$$

25 MatchedTemplatePeaks = Normalized Areas of the matching template peaks  
 MatchedTestPeaks = Normalized Areas of the matching test peaks

In the case where the alignment position of the three consecutive peaks is the same for both the template and the test  
 30 IVEGs (i.e., the test and template complexes match), Equation (3) reduces to Equation (4):

Equation (4)

35 
$$\text{Score} = |\text{TemplatePeak}_1 - \text{TestPeak}_1| + |\text{TemplatePeak}_2 - \text{TestPeak}_2| + |\text{TemplatePeak}_3 - \text{TestPeak}_3|$$

where:

40 TemplatePeak<sub>1</sub> = Normalized Area of dominant template peak  
 TestPeak<sub>1</sub> = Normalized Area of dominant test peak  
 TemplatePeak<sub>2</sub> = Normalized Area of second dominant template peak  
 TestPeak<sub>2</sub> = Normalized Area of second dominant test peak  
 TemplatePeak<sub>3</sub> = Normalized Area of third dominant template peak  
 45 TestPeak<sub>3</sub> = Normalized Area of third dominant test peak

Thus, in the preferred embodiment, the method uses Equation (3) when some peaks have no match in the test and template IVEGs and Equation (4) when a match is found between all three peaks of the test and template IVEGs. If desired, however, Equation (3) may be used in all cases.

50 [0054] As can be seen from Equations (3) and (4), the score indicates the relative similarity between the test and template IVEGs. In the preferred embodiment, a smaller score indicates relative similarity, while a larger score indicates that the two IVEGs are dissimilar.

[0055] FIG. 8 illustrates a system 802 according to a preferred embodiment of the present invention. Although applicable to signals in general, the system illustrated in FIG. 8 is directed toward the preferred embodiment of IVEGs in an implantable cardiac pacing or defibrillation device. The system contains a template collection unit 804, a test IVEG collection unit 808, a peak information extraction unit 810, an alignment unit 812, a score determination unit 814, and a threshold comparison unit 816. In addition, the system includes a template storage area 806.

[0056] The template collection unit 804 creates a template as described above. In the preferred embodiment the template so created is stored in a template storage area 806. Thus, a template need not be created each time a test IVEG is to be compared to a template. Rather a template can be retrieved from the template storage area 806 and passed to that template collection unit 804 for subsequent processing in the system.

5 [0057] The test IVEG collection unit 808 collects a test IVEG for determination of whether the test IVEG is a sinus or VT complex. The test IVEG collection unit 808 collects a test IVEG during a sense refractory period in a well-known manner. The test IVEG collection unit 808 and template collection unit 804 are coupled to a peak information extraction unit 810.

10 [0058] The peak information extraction unit 810 extracts peak information of interest from both the template and test IVEGs for the comparison phase of the present invention. As explained above, the peak information of interest according to a preferred embodiment are the normalized area, polarity, and position of each of the three consecutive peaks having the largest sum of peak amplitudes in a collected signal. In addition, the peak information extraction unit 810 normalizes the areas of the 3 consecutive peaks. The peak information extraction unit 810 is coupled to an alignment unit 812.

15 [0059] The alignment unit 812 determines whether the template and test IVEGs are aligned as described above. The alignment unit 812 is coupled to a score determination unit 814. The alignment unit 812 transmits an indication of whether the test and template signals match to the score determination unit 814.

20 [0060] The score determination unit 814 receives the match indication from the alignment unit 812. The score determination unit calculates a score using Equation (3) or Equation (4), as described above. Whether the system uses Equation (3) or Equation (4), depends on the received match indication. If the received match indication indicates no match, the system uses Equation (3). If on the other hand, the received match indication indicates a match, the system uses Equation (4). The score determination unit is coupled to a threshold comparison unit 816. Note that the system, if desired, could use Equation (3) regardless of whether a match is indicated.

25 [0061] The threshold comparison unit 816 compares the score to a threshold (described above). If the score is greater than the threshold, then the test and template signals are dissimilar. However, if the score is less than the threshold, the test and template signals are similar to one another.

30 [0062] FIG. 9 illustrates the preferred embodiment for the system described in FIG. 8. The preferred embodiment includes a microprocessor 902, a random access memory (RAM) 904, an input/output (I/O) unit 908, and a bus 906. The functionality of system 802 is implemented in software in microprocessor 902. Bus 906 provides for communication between RAM 904, microprocessor 902 and input/output unit 908. I/O unit 908 is also coupled to a patient's heart 910 in a well known manner to collect the template and test IVEGs.

35 [0063] In FIG. 9, microprocessor 902 instructs I/O unit 908 to collect a template IVEG. I/O unit 908 collects the template IVEG in a well-known manner. I/O unit 908 stores the template IVEG in RAM 904 over bus 906. Microprocessor 902 extracts peak information from the template IVEG and stores the peak information in RAM 904. The original template IVEG can be discarded. At a later time, microprocessor 902 instructs I/O unit 908 to collect a test IVEG. The test IVEG is stored in RAM 904 over bus 906. Microprocessor 902 extracts peak information from the test IVEG and stores the information in RAM 904. The original test IVEG can then be discarded. After obtaining the peak information from the test IVEG, microprocessor 902 performs the alignment of alignment unit 812 and determines the score as explained with reference to score determination unit 814. Microprocessor 902 then performs a threshold comparison test as explained with reference to threshold comparison unit 816 to determine whether the test and template IVEGs match.

40 [0064] While various embodiments of the present invention have been described above, it should be understood that they have been presented by way of example only, and not limitation. Thus, the breadth and scope of the present invention should not be limited by any of the above-described exemplary embodiments, but should be defined only in accordance with the following claims.

45

## Claims

- 50 1. An apparatus for distinguishing between a ventricular tachycardia (VT) complex and a sinus tachycardia (ST) or supra ventricular tachycardia (SVT) complex, comprising:
  - (a) means (908; 804, 808) for collecting a template intraventricular electrogram (700) and a test intraventricular electrogram (710);
  - (b) means (902; 303) for locating a group of at least two consecutive peaks (210, 212, 214) having a largest cumulative area for each of said intraventricular electrograms (700, 710),
  - (c) means (902; 306, 308) for determining a polarity and a position of each peak (210, 212, 214) within said group for each of said intraventricular electrograms (700, 710),
  - (d) means (902; 310) for normalizing the area of each peak (210, 212, 214) in said group, wherein said polar-

- 5                    ities, said positions and said normalized areas represent extracted peak information;  
 (e) means (902; 504, 510; 604) for comparing polarities and positions of said extracted peak information for  
 said test intraventricular electrogram (710) with polarities and positions of said extracted information for said  
 template intraventricular electrogram (700) to align said test intraventricular electrogram (710) with said tem-  
 plate intraventricular electrogram (700); and  
 (f) means (902; 111) for comparing said extracted peak information for said test intraventricular electrogram  
 (710) with said extracted information for said template intraventricular electrogram (700) to generate a score  
 indicative of similarity between said template intraventricular electrogram (700) and said test intraventricular  
 electrogram (710).
- 10                 2. The apparatus of claim 1, **characterized in that** said means (902; 310) for normalizing comprises:  
 means (902; 310) for normalizing each peak (210, 212, 214) with respect to the sum of the areas of each of  
 said three consecutive peaks (210, 212, 214).
- 15                 3. The apparatus of claim 1, **characterized in that** said means (902; 310) for normalizing comprises:  
 means (902; 310) for normalizing each peak (210, 212, 214) with respect to a peak (212) having the largest  
 area of said three consecutive peaks (210, 212, 214).
- 20                 4. The apparatus of any of claims 1 - 3, **characterized in that** said means (902; 406; 504, 510; 604) for comparing  
 includes:  
 means (504) for comparing the polarity of a dominant test peak to the polarity of a dominant template peak;  
 means (406) for aligning said template intraventricular electrogram (700) with said test intraventricular elec-  
 trogram (710) on the basis of said dominant peaks if the polarity of said dominant test peak matches the  
 polarity of said dominant template peak;  
 means (510) for comparing the polarity of said dominant test peak to the polarity of said second dominant  
 template peak if the polarity of said dominant test peak does not match the polarity of said dominant template  
 peak;  
 means (406) for aligning said template intraventricular electrogram (700) with said test intraventricular elec-  
 trogram (710) on the basis of said dominant test peak and said second dominant template peak if the polarity  
 of said dominant test peak matches the polarity of said second dominant template peak; and  
 means (114) for indicating that the test and template intraventricular electrograms (700, 710) could not be  
 aligned if the polarity of said dominant test peak does not match the polarity of either said dominant template  
 peak or said second dominant template peak.
- 25                 5. The apparatus of claim 4, **characterized in that** said means (406) for aligning includes:  
 means (604) for determining whether positions of said test peaks correspond to positions of said template  
 peaks; and  
 means (605, 606) for scanning said test intraventricular electrogram (710) for another peak with which a match  
 can be made if said positions of said test peaks do not correspond to positions of said template peaks.
- 30                 45                 6. The apparatus of any of claims 1 - 5, **characterized in that** said means (902; 111) for comparing said extracted  
 peak information comprises:  
 means for calculating said score as the sum of the magnitudes of the areas of unmatched template peaks,  
 the areas of unmatched test peaks, and the difference between the areas of any matched template and test  
 peaks if peaks in said template intraventricular electrogram (700) and said test intraventricular electrogram  
 (710) do not match in position and polarity; and  
 means for calculating said score as the sum of the magnitudes of the differences between the areas of the  
 matched peaks if peaks in said template intraventricular electrogram (700) and said test intraventricular elec-  
 trogram (710) all match in position and polarity.
- 35                 55                 7. A method for distinguishing between a ventricular tachycardia complex and a sinus tachycardia or supra ventricular  
 tachycardia complex, comprising the steps of:

- (a) collecting a template intraventricular electrogram and a test intraventricular electrogram;  
5 (b) for each of said intraventricular electrograms,  
      (1) locating a group of at least two consecutive peaks having a largest cumulative area,  
      (2) determining a polarity and a position of each peak within said group, and  
      (3) normalizing the area of each peak in said group, wherein said polarities, said positions and said normalized areas represent extracted peak information;
- 10 (c) comparing polarities and positions of said extracted peak information for said test intraventricular electrogram with polarities and positions of said extracted information for said template intraventricular electrogram to align said test intraventricular electrogram with said template intraventricular electrogram; and  
      (d) comparing said extracted peak information for said test intraventricular electrogram with said extracted information for said template intraventricular electrogram to generate a score indicative of similarity between said template intraventricular electrogram and said test intraventricular electrogram.
- 15

**Patentansprüche**

- 20 1. Vorrichtung zum Unterscheiden zwischen einem ventrikulären Tachykardie- (VT) -Komplex und einem Sinus-Tachykardie- (ST) oder supra-ventrikulärem Tachykardie- (SVT) -Komplex, mit:  
      (a) Mitteln (908; 804; 808) zum Aufnehmen eines intraventrikulären Schablonen-Elektrogramms (700) und eines intraventrikulären Test-Elektrogramms (710);  
      (b) Mitteln (902; 303) zum Auffinden einer Gruppe von wenigstens zwei aufeinanderfolgenden Spitzenwerten (210, 212, 214), die eine größte kumulative Fläche besitzt, und zwar für jedes der intraventrikulären Elektrogramme (700, 710);  
      (c) Mitteln (902; 306; 308) zum Bestimmen einer Polarität und einer Position von jedem Spitzenwert (210, 212, 214) innerhalb der Gruppe, und zwar für jedes der intraventrikulären Elektrogramme (700, 710);  
      (d) Mitteln (902; 310) zum Normieren der Fläche von jedem Spitzenwert (210, 212, 214) in der Gruppe, wobei die Polaritäten, die Positionen und die normierten Flächen extrahierte Spitzenwertinformation darstellen;  
      (e) Mitteln (902; 504; 510; 604) zum Vergleichen von Polaritäten und Positionen der extrahierten Spitzenwertinformation für das intraventrikuläre Test-Elektrogramm (710) mit Polaritäten und Positionen der extrahierten Information für das intraventrikuläre Schablonen-Elektrogramm (700), und zwar um das intraventrikuläre Test-Elektrogramm (710) mit dem intraventrikulären Schablonen-Elektrogramm (700) auszurichten; und  
      (f) Mitteln (902; 111) zum Vergleichen der extrahierten Spitzenwertinformation für das intraventrikuläre Test-Elektrogramm (710) mit der extrahierten Information für das intraventrikuläre Schablonen-Elektrogramm (700), und zwar um ein Ergebnis zu erzeugen, das die Ähnlichkeit zwischen dem intraventrikulären Schablonen-Elektrogramm (700) und dem intraventrikulären Test-Elektrogramm (710) anzeigt.
- 25 40 2. Vorrichtung nach Anspruch 1, dadurch gekennzeichnet, dass die Mittel (902; 310) zum Normieren aufweisen:  
      Mittel (902; 310) zum Normieren eines jeden Spitzenwertes (210, 212, 214) in Bezug auf die Summe der Flächen der drei aufeinander folgenden Spitzenwerte (210, 212, 214).
- 30 45 3. Vorrichtung nach Anspruch 1, dadurch gekennzeichnet, dass die Mittel (902; 310) zum Normieren aufweisen:  
      Mittel (902; 310) zum Normieren von jedem Spitzenwert (210, 212, 214) in Bezug auf einen Spitzenwert (212), der die größte Fläche von den drei aufeinander folgenden Spitzenwerten (210, 212, 214) besitzt.
- 35 50 55 4. Vorrichtung nach einem der Ansprüche 1 bis 3, dadurch gekennzeichnet, dass die Mittel (902; 406; 504; 510; 604) zum Vergleichen aufweisen:  
      Mittel (504) zum Vergleichen der Polarität eines dominanten Test-Spitzenwertes mit der Polarität eines dominanten Schablonen-Spitzenwertes;  
      Mittel (406) zum Ausrichten des intraventrikulären Schablonen-Elektrogramms (700) mit dem intraventrikulären Test-Elektrogramm (710), und zwar auf der Grundlage der dominanten Spitzenwerte, dann, wenn die Polarität des dominanten Test-Spitzenwertes mit der Polarität des dominanten Schablonen-Spitzenwertes übereinstimmt;

- Mittel (510) zum Vergleichen der Polarität des dominanten Test-Spitzenwertes mit der Polarität des zweiten dominanten Schablonen-Spitzenwertes, dann, wenn die Polarität des dominanten Test-Spitzenwertes nicht mit der Polarität des dominanten Schablonen-Spitzenwertes übereinstimmt;
- 5 Mittel (406) zum Ausrichten des intraventrikulären Schablonen-Elektrogramms (700) mit dem intraventrikulären Test-Elektrogramm (710) auf der Grundlage des dominanten Test-Spitzenwertes und des zweiten dominanten Schablonen-Spitzenwertes, dann, wenn die Polarität des dominanten Testspitzenwertes mit der Polarität des zweiten dominanten Schablonen-Spitzenwertes übereinstimmt; und
- 10 Mittel (114) zum Anzeigen, dass das intraventrikuläre Test und das intraventrikuläre Schablonen-Elektrogramm (700, 710) nicht ausgerichtet werden konnten, und zwar dann, wenn die Polarität des dominanten Test-Spitzenwertes weder mit der Polarität des dominanten Schablonen-Spitzenwertes noch mit jener des zweiten dominanten Schablonen-Spitzenwertes übereinstimmt.
5. Vorrichtung nach Anspruch 4, dadurch gekennzeichnet, dass die Mittel (406) zum Ausrichten aufweisen:
- 15 Mittel (604) zum Bestimmen, ob Positionen der Test-Spitzenwerte Positionen der Schablonen-Spitzenwerte entsprechen; und
- Mittel (605, 606) zum Scannen des intraventrikulären Test-Elektrogramms (710) nach einem weiteren Spitzenwert, mit dem eine Übereinstimmung vorgenommen werden kann, wenn die Positionen der Test-Spitzenwerte nicht Positionen der Schablonen-Spitzenwerte entsprechen.
- 20
6. Vorrichtung nach einem der Ansprüche 1 bis 5, dadurch gekennzeichnet, dass die Mittel (902; 111) zum Vergleichen der extrahierten Spitzenwertinformation aufweisen:
- 25 Mittel zum Berechnen des Ergebnisses als die Summe der Größen der Flächen von nicht übereinstimmenden Schablonen-Spitzenwerten, der Flächen der nicht übereinstimmenden Test-Spitzenwerte und der Differenz zwischen den Flächen von irgendwelchen übereinstimmenden Schablonen- und Test-Spitzenwerten, und zwar dann, wenn Spitzenwerte in dem intraventrikulären Schablonen-Elektrogramm (700) und dem intraventrikulären Test-Elektrogramm (710) hinsichtlich Position und Polarität nicht übereinstimmen; und
- 30 Mittel zum Berechnen des Ergebnisses als die Summe der Größen der Differenzen zwischen den Flächen der übereinstimmenden Spitzenwerte, und zwar dann, wenn Spitzenwerte in dem intraventrikulären Schablonen-Elektrogramm (700) und dem intraventrikulären Test-Elektrogramm (710) sämtlich hinsichtlich Position und Polarität übereinstimmen.
- 35
7. Verfahren zum Unterscheiden zwischen einem ventrikulären Tachykardie-Komplex und einem Sinus-Tachykardie- oder supraventrikulären Tachykardie-Komplex, mit den Schritten:
- 40
- a) Aufnehmen eines intraventrikulären Schablonen-Elektrogramms und eines intraventrikulären Test-Elektrogramms;
- (b) für jedes der intraventrikulären Elektrogramme
- 45
- (1) Auffinden einer Gruppe von wenigstens zwei aufeinander folgenden Spitzenwerten, die eine größte kumulative Fläche besitzen,
- (2) Bestimmen einer Polarität und einer Position von jedem Spitzenwert innerhalb der Gruppe, und
- 50
- (3) Normieren der Fläche von jedem Spitzenwert in der Gruppe, wobei die Polaritäten, die Positionen und die normierten Flächen extrahierte Spitzenwertinformation darstellen;
- 55
- (c) Vergleichen von Polaritäten und von Positionen der extrahierten Spitzenwertinformation für das intraventrikuläre Test-Elektrogramm mit Polaritäten und Positionen der extrahierten Information für das intraventrikuläre Schablonen-Elektrogramm, um das intraventrikuläre Test-Elektrogramm mit dem intraventrikulären Schablonen-Elektrogramm auszurichten; und
- (d) Vergleichen der extrahierten Spitzenwertinformation für das intraventrikuläre Test-Elektrogramm mit der extrahierten Information für das intraventrikuläre Schablonen-Elektrogramm, um ein Ergebnis zu erzeugen, das die Ähnlichkeit zwischen dem intraventrikulären Schablonen-Elektrogramm und dem intraventrikulären Test-Elektrogramm anzeigt.

**Revendications**

1. Appareil permettant de faire la distinction entre un complexe de tachycardie ventriculaire (TV) et un complexe de tachycardie sinusal (TS) ou supraventriculaire (TSV), comprenant :

5

(a) un moyen (908 ; 804 ; 808) permettant de collecter un électrogramme intraventriculaire modèle (700) et un électrogramme intraventriculaire test (710) ;  
 (b) un moyen (902 ; 303) permettant de localiser un groupe d'au moins deux pics consécutifs (210, 212, 214) possédant la zone cumulative la plus large pour chacun desdits électrogrammes intraventriculaires (700, 710),  
 10 (c) un moyen (902 ; 306 ; 308) permettant de déterminer la polarité et la position de chaque pic (210, 212, 214) à l'intérieur dudit groupe pour chacun desdits électrogrammes intraventriculaires (700, 710) ;  
 (d) un moyen (902 ; 310) permettant de normaliser la zone de chaque pic (210, 212, 214) dans ledit groupe, dans lequel lesdites polarités, lesdites positions et lesdites zones normalisées représentent des informations de pic extraites ;  
 15 (e) un moyen (902 ; 504 ; 510 ; 604) permettant de comparer les polarités et les positions desdites informations de pic extraites pour ledit électrogramme intraventriculaire test (710) avec les polarités et les positions desdites informations extraites pour ledit électrogramme intraventriculaire modèle (700) afin d'aligner ledit électrogramme intraventriculaire test (710) et ledit électrogramme intraventriculaire modèle (700) ; et  
 20 (f) un moyen (902 ; 111) permettant de comparer lesdites informations de pic extraites pour ledit électrogramme intraventriculaire test (710) avec lesdites informations extraites pour ledit électrogramme intraventriculaire modèle (700) afin de générer un indice indicatif des similarités entre ledit électrogramme intraventriculaire modèle (700) et ledit électrogramme intraventriculaire test (710).

2. Appareil selon la revendication 1, caractérisé en ce que ledit moyen (902 ; 310) de normalisation comprend :

25

un moyen (902 ; 310) permettant de normaliser chaque pic (210, 212, 214) par rapport à la somme des zones de chacun desdits trois pics consécutifs (210, 212, 214).

3. Appareil selon la revendication 1, caractérisé en ce que ledit moyen (902 ; 310) de normalisation comprend :

30

un moyen (902 ; 310) permettant de normaliser chaque pic (210, 212, 214) par rapport à un pic (212) possédant la zone la plus large desdits trois pics consécutifs (210, 212, 214).

4. Appareil selon l'une quelconque des revendications 1 à 3, caractérisé en ce que ledit moyen (902 ; 406 ; 504 ; 510 ; 604) de comparaison comprend :

35

un moyen (504) permettant de comparer la polarité d'un pic dominant du test à la polarité d'un pic dominant du modèle ;

un moyen (406) permettant d'aligner ledit électrogramme intraventriculaire modèle (700) au dit électrogramme intraventriculaire test (710) sur la base desdits pics dominants si la polarité dudit pic dominant du test correspond à la polarité dudit pic dominant du modèle ;

un moyen (510) permettant de comparer la polarité dudit pic dominant du test à la polarité dudit second pic dominant du modèle si la polarité dudit pic dominant du test ne correspond pas à la polarité dudit pic dominant du modèle ;

un moyen (406) permettant d'aligner ledit électrogramme intraventriculaire modèle (700) avec ledit électrogramme intraventriculaire test (710) sur la base dudit pic dominant du test et dudit second pic dominant du modèle si la polarité dudit pic dominant du test correspond à la polarité dudit second pic dominant du modèle ; et

un moyen (114) permettant d'indiquer que lesdits électrogrammes intraventriculaires test et modèle (700 et 710) ne peuvent pas être alignés si la polarité dudit pic dominant du test ne correspond pas à la polarité soit dudit pic dominant du modèle, soit dudit second pic dominant du modèle.

45

5. Appareil selon la revendication 4, caractérisé en ce que ledit moyen (406) d'alignement comprend :

50

un moyen (604) permettant de déterminer si les positions desdits pics du test correspondent aux positions desdits pics du modèle ; et

un moyen (605, 606) permettant de balayer ledit électrogramme intraventriculaire test (710) pour trouver un autre pic avec lequel il soit possible d'établir une correspondance si lesdites positions desdits pics du test ne correspondent pas aux positions desdits pics du modèle.

6. Appareil selon l'une quelconque des revendications 1 à 5, caractérisé en ce que ledit moyen (902 ; 111) de comparaison desdites informations de pic extraites comprend :

5        un moyen permettant de calculer ledit indice comme la somme des amplitudes des zones des pics sans correspondance du modèle, des zones des pics sans correspondance du test, et de la différence entre les zones des pics correspondants du modèle et du test si la position et la polarité des pics dudit électrogramme intraventriculaire modèle (700) et dudit électrogramme intraventriculaire test (710) ne correspondent pas ; et  
10      un moyen permettant de calculer ledit indice comme la somme des amplitudes de la différence entre les zones des pics correspondants si la position et la polarité des pics dudit électrogramme intraventriculaire modèle (700) et dudit électrogramme intraventriculaire test (710) correspondent toutes.

7. Procédé permettant de faire la distinction entre un complexe de tachycardie ventriculaire et un complexe de tachycardie sinusale ou supraventriculaire, comprenant les étapes consistant à :

15        (a) collecter un électrogramme intraventriculaire modèle et un électrogramme intraventriculaire test ;  
              (b) pour chacun desdits électrogrammes intraventriculaires,  
                        (1) localiser un groupe d'au moins deux pics consécutifs ayant la zone cumulative la plus large,  
                        (2) déterminer la polarité et la position de chaque pic à l'intérieur dudit groupe,  
20        et  
                        (3) normaliser la zone de chaque pic dans ledit groupe, dans lequel lesdites polarités, lesdites positions et lesdites zones normalisées représentent des informations de pic extraites ;  
              (c) comparer les polarités et les positions desdites informations de pic extraites pour ledit électrogramme intraventriculaire test avec les polarités et les positions desdites informations extraites pour ledit électrogramme intraventriculaire modèle afin d'aligner ledit électrogramme intraventriculaire test avec ledit électrogramme intraventriculaire modèle ; et  
              (d) comparer lesdites informations de pic extraites pour ledit électrogramme intraventriculaire test avec lesdites informations extraites pour ledit électrogramme intraventriculaire modèle afin de générer un indice indicatif des similarités entre ledit électrogramme intraventriculaire modèle et ledit électrogramme intraventriculaire test.  
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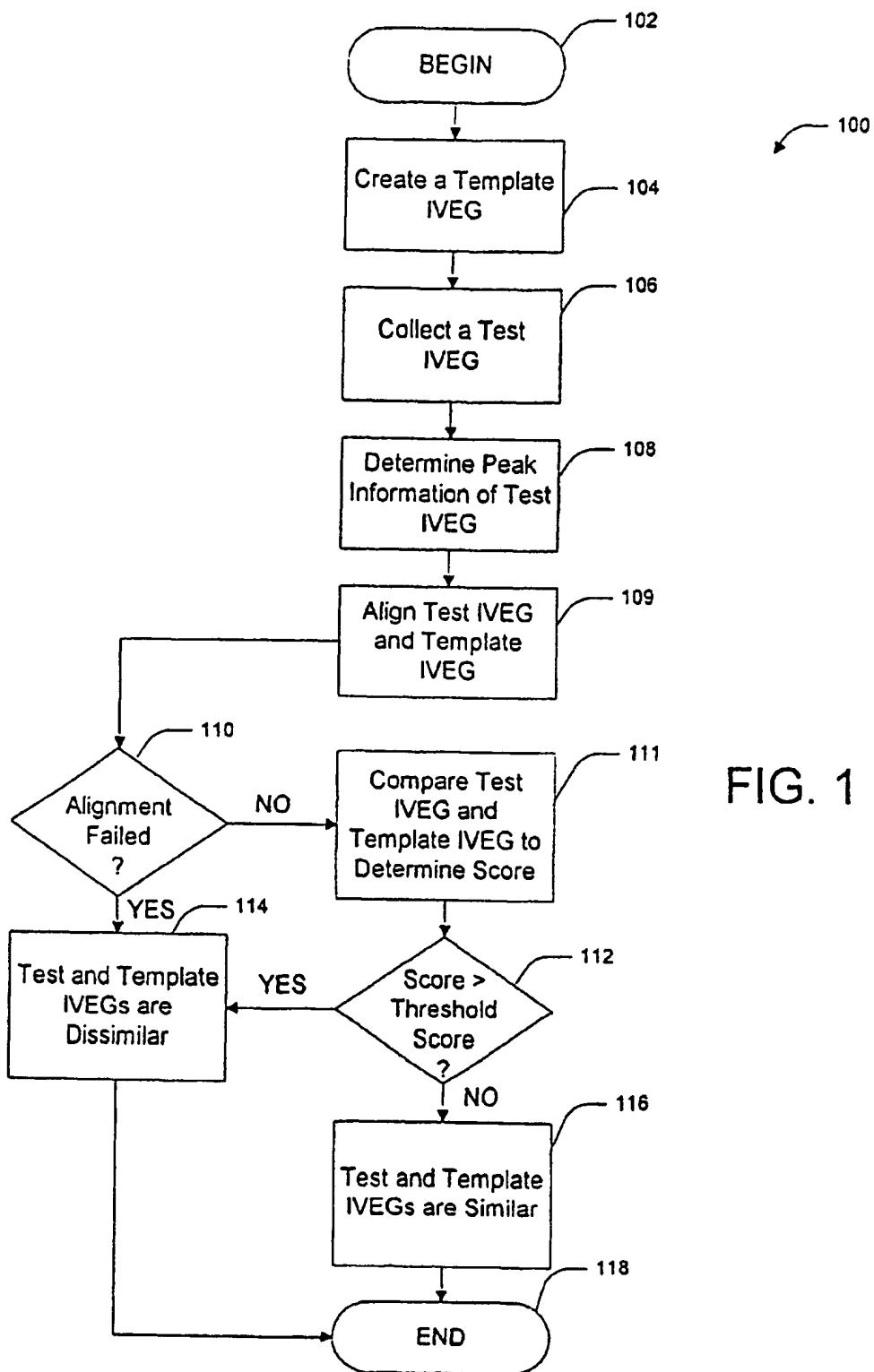


FIG. 1

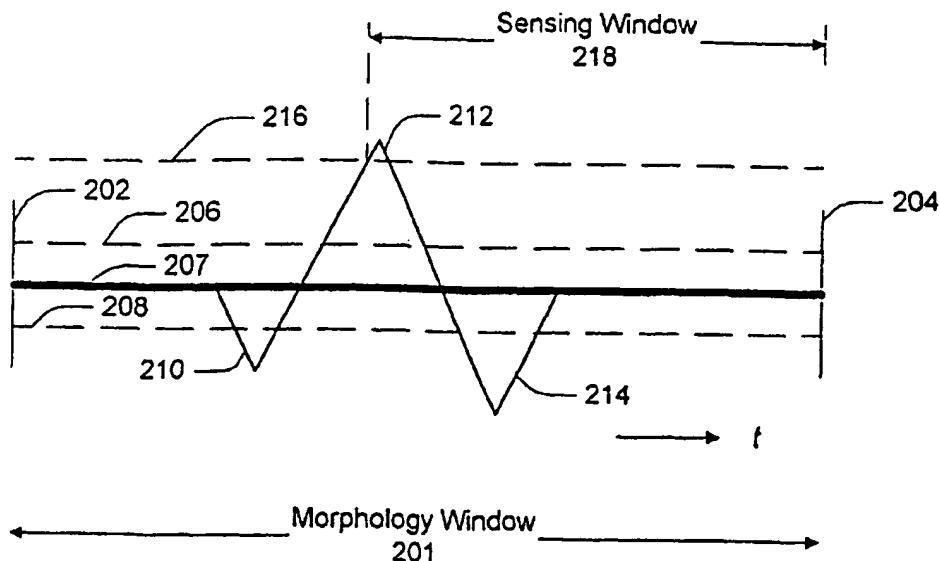


FIG. 2

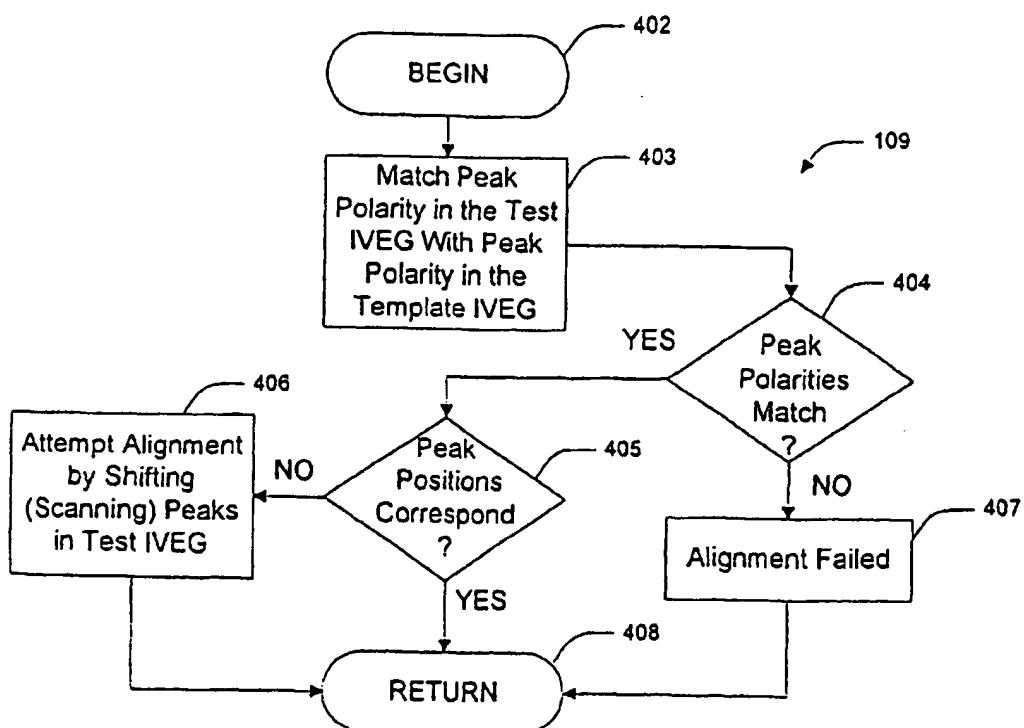


FIG. 4

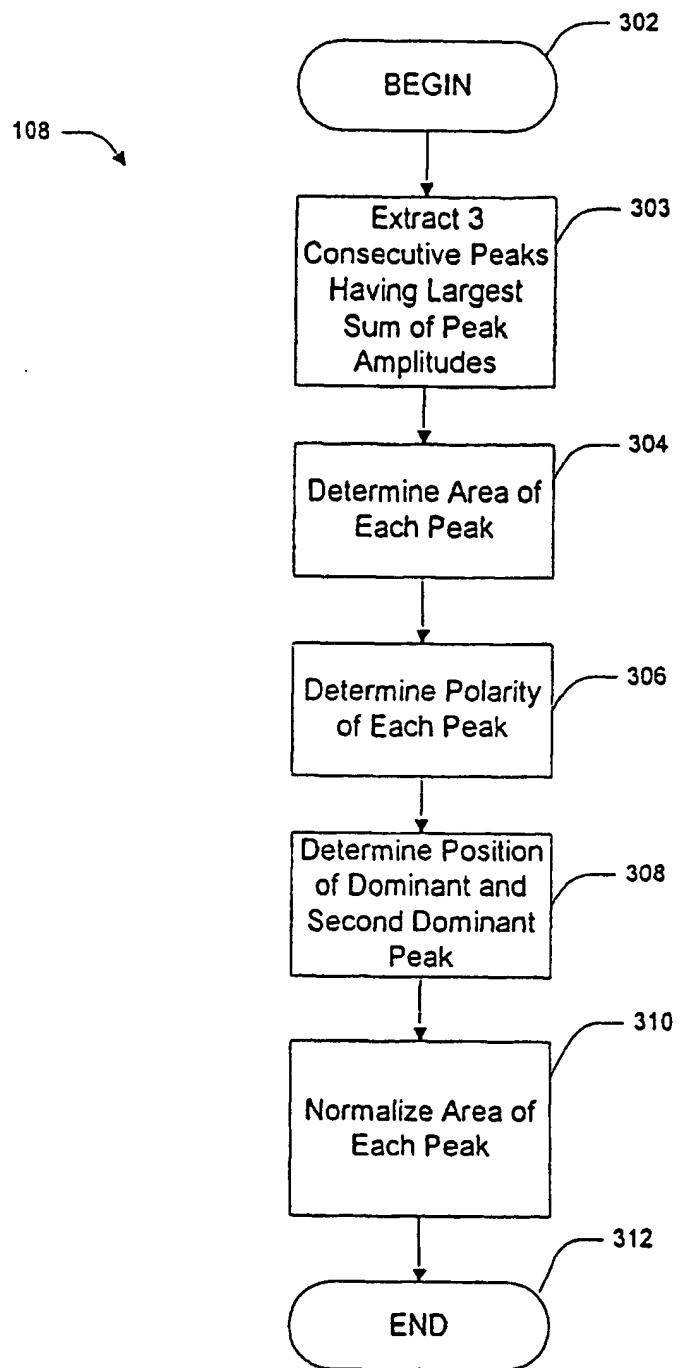


FIG. 3

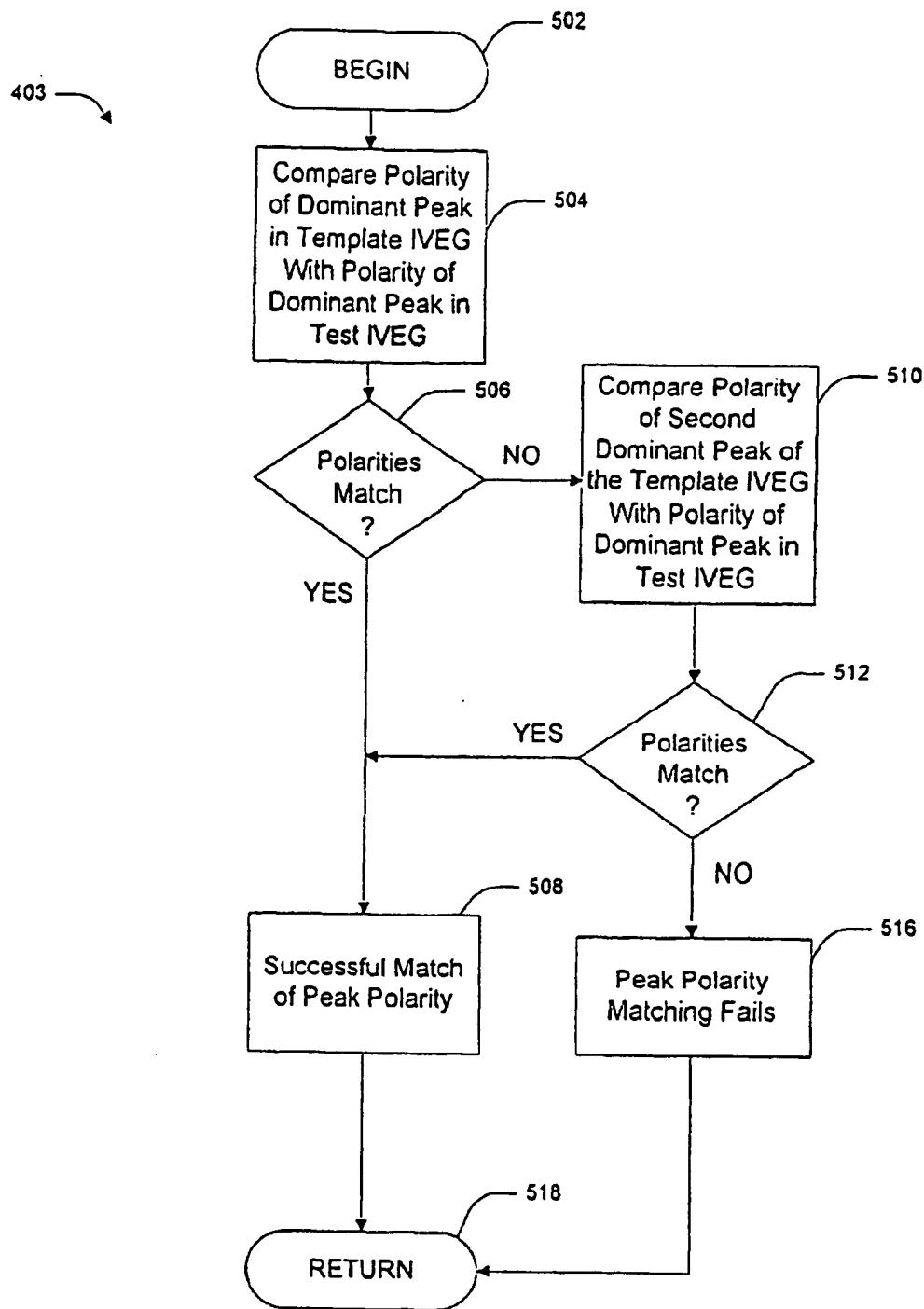


FIG. 5

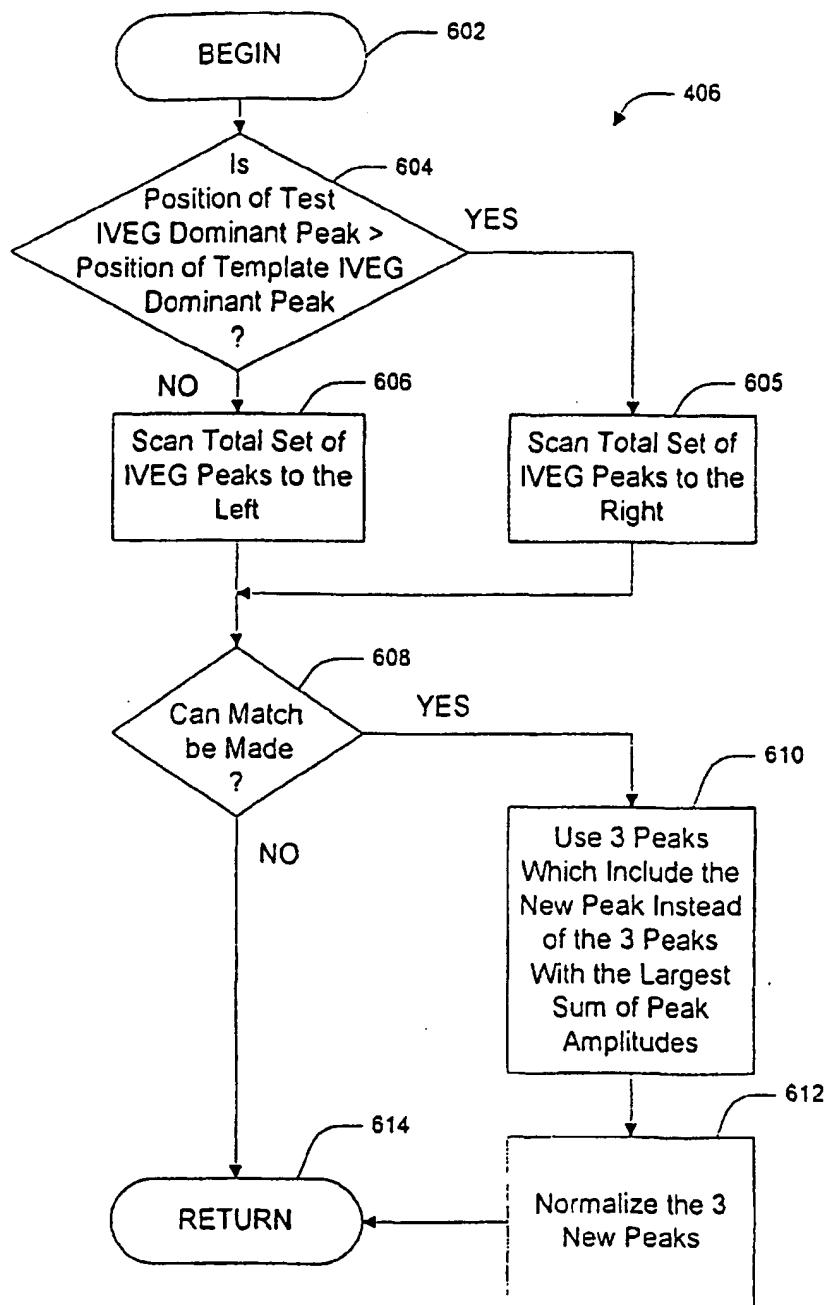


FIG. 6

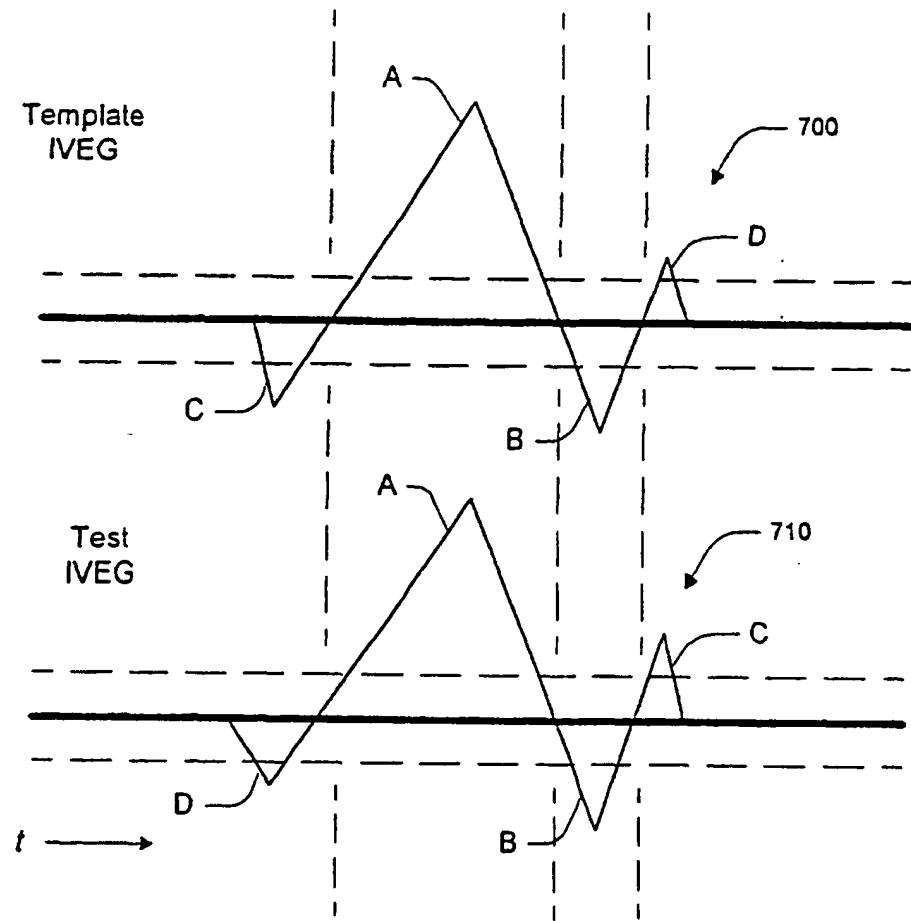


FIG. 7

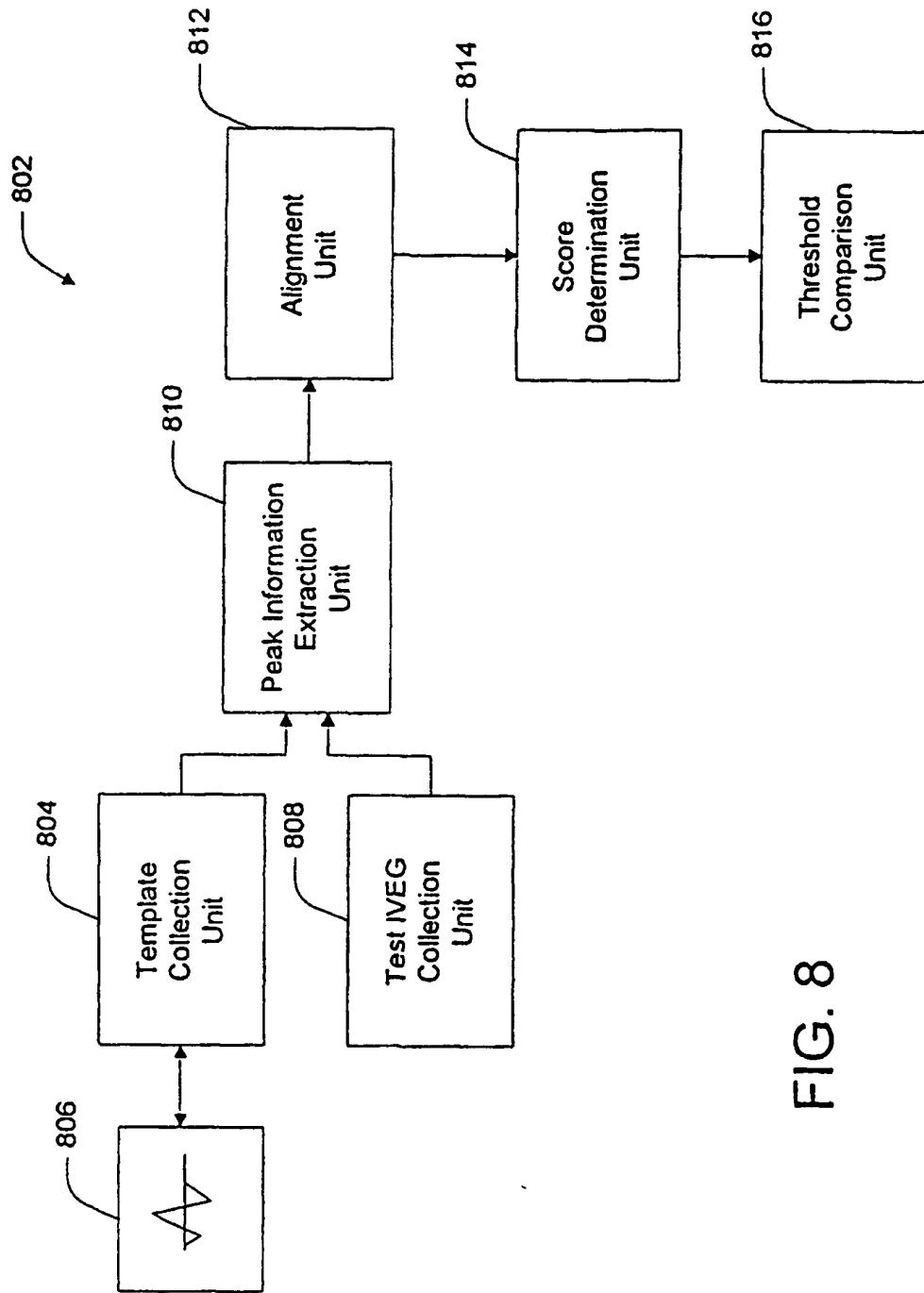


FIG. 8

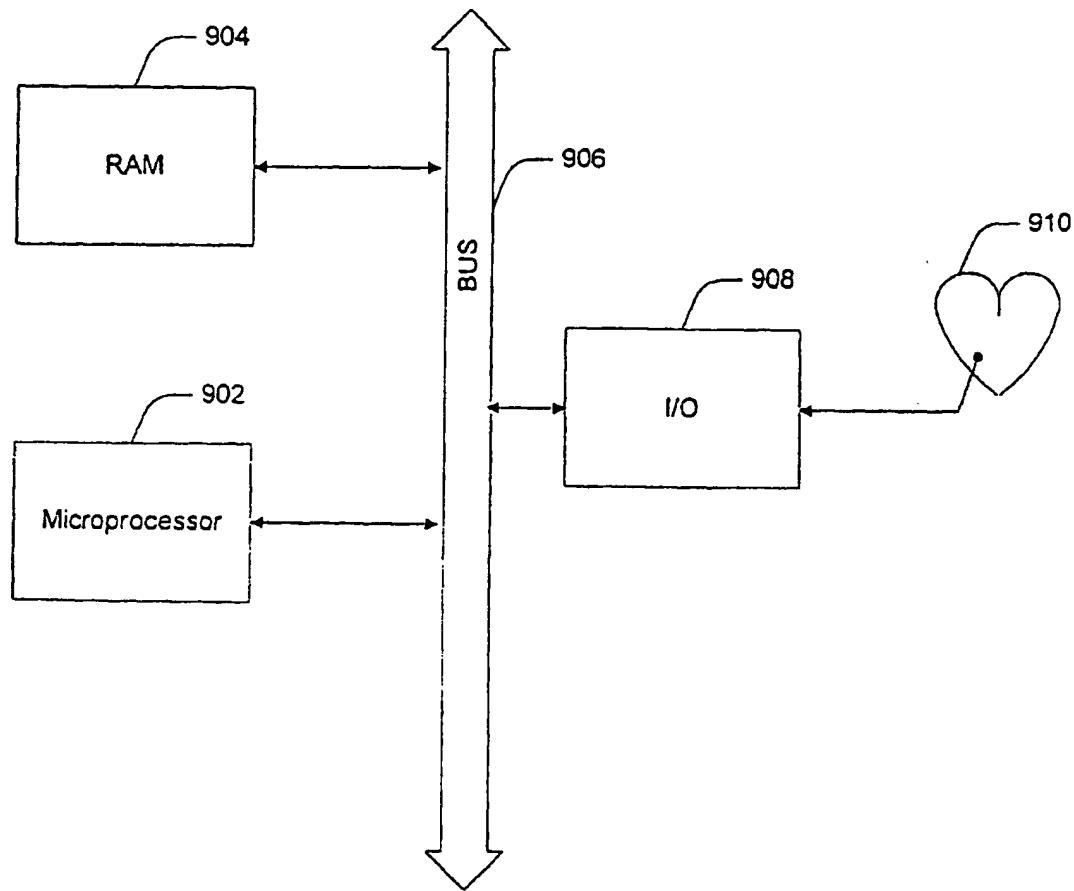


FIG. 9